- (6) a Supplemental Information Disclosure Statement (in duplicate), accompanied by the appropriate provision authorizing payment of the required fee;
 - (7) a revised PTO 1449 form and copies of the references listed therein; and
 - (8) a Request for Continued Examination with the appropriate fees.

IN THE CLAIMS:

Please amend the claims as follows:

Cancel Claims 25-27 without prejudice.

Amend Claims 5, 6, 16, and 28-30 as follows.

Add new Claim 31.

The pending claims read as follows:

- 2. (Previously Amended) The method of Claim 28, wherein the needle is selected from the group consisting of microneedles, catheter needles, and injection needles.
- 3. (Previously Amended) The method of Claim 28, wherein a single needle is inserted.
- 4. (Previously Amended) The method of Claim 28, wherein multiple needles are inserted.
- 5. (Currently Amended) The method of any of Claims 25-30 28, 29, or 31, wherein the substance is a liquid delivered by pressure directly on the liquid.
- 6. (Currently Amended) The method of any of Claims 25-30 28, 29, or 31, wherein a hormone is delivered.
- 7. (Previously Amended) The method of Claim 6, wherein the hormone is selected from the group consisting of insulin and PTH.
- 10. (Previously Amended) The method of Claim 28, wherein the needle is about 300 mm to 2 mm long.
- 11. (Previously Amended) The method of Claim 28, wherein the needle is about 500 mm to 1 mm long.
- 12. (Previously Amended) The method of Claim 28, wherein the outlet is at a depth of about 250 mm to 2 mm when the needle is inserted.
- 13. (Previously Amended) The method of Claim 28, wherein the outlet is at a depth of about 750 mm to 1.5 mm when the needle is inserted.

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- 14. (Previously Amended) The method of Claim 28, wherein the outlet has an exposed height of about 0 to 1 mm.
- 15. (Previously Amended) The method of Claim 28, wherein the outlet has an exposed height of about 0 to 300 mm.
- 16. (Currently Amended) The method of Claim 26 28, wherein the delivery rate or volume is controlled by spacing of multiple needles, needle diameter or number of needles.
- 25. (Canceled)
- 26. (Canceled)
- 27. (Canceled)
- 28. (Currently Amended) A method for the administration of a substance to a human subject, comprising delivering the substance into the an intradermal compartment of the human subject's skin at a controlled volume and rate via a needle having a length sufficient to penetrate the intradermal space and an outlet at a depth within the intradermal space so that the substance is distributed systemically in the plasma.
- 29. (Currently Amended) A method for the administration of a substance to a human subject, comprising delivering the substance into the <u>an</u> intradermal compartment of the human subject's skin, so that the substance is distributed systemically and has a pharmacokinetic profile similar to subcutaneous delivery of the substance, but with <u>a</u> higher plasma levels.
- 30. (Currently Amended) A method for the administration of a substance to a human subject, comprising delivering the substance into the <u>an</u> intradermal compartment of the human subject's skin, so that the substance is distributed systemically and has a pharmacokinetic profile similar to subcutaneous delivery of the substance, but with a faster onset of <u>a</u> detectable plasma levels.
- 31. (New) A method for the administration of a substance to a human subject, comprising delivering the substance into the an intradermal compartment of the human subject's skin, so that the substance is distributed systemically and has a pharmacokinetic profile similar to subcutaneous delivery of the substance, but with a higher bioavailability.

REMARKS

First and foremost, Applicants would like to thank Examiners Tran, Denion, Hayes, and Casler for their cooperation in organizing an interview with the Applicants on May 14,